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Substitute for Form 1449A/PTO INFORMATION DISCLOSURE STATEMENT BY APPLICANT				Application Number		
				Filing Date		Herewith
				First Named Inventor		Takahide NISHI et al.
				Group Art Unit		
				Examiner Name		
Sheet	1	of	3	Attorney Docket Number		02697CIPD/HG

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Exam. Inits ¹	Cite No ²	Document Number	Kind Code ³	Name of Patentee or Applicant	Publication Date MM-DD-YYYY	Relevant Portion
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Examiner Signature		<i>D. L. [Signature]</i>			Date Considered	<i>3/2/05</i>		

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Examiner Initials ¹	Cite No. ¹	Include name of author (in CAPITAL LETTERS), title of article, title of item, date, page(s), volume-issue number(s), publisher, city and/or country where published					T ²
<i>DM</i>		<p>HUANG, et al., Kidney International, "Th1 responsiveness to nephritogenic antigens determines susceptibility to crescentic glomerulonephritis in mice", <u>51</u>, pp. 94-103 (1997)</p> <p>BERMAN, et al., J. Immunology, "Decreased Il-4 Production in New Onset Type I Insulin-Dependent Diabetes Mellitus¹", <u>157</u>, pp. 4690-4696 (1996)</p> <p>CATIVIELA, et al., "Stereoselective synthesis of quaternary α-amino acids. Part 1: Acyclic compounds", Tetrahedron: Asymmetry, <u>2</u>, pp. 3517-3599 (1998)</p> <p>GANDER, et al., "Synthesis of Enantiomerically Pure, α-Alkylated Lysine, Ornithine, and Tryptophan-Derivatives", Helvetica Chimica Acta, <u>71</u>, pp. 224-236, (1988) (English language Abstract)</p> <p>SANO, et al., "Lewis Acid- and Cationic Lithium-Mediated Diastereoselective Aldol-Type Reaction Based on a Double Chiral Recognition manner for the Asymmetric Synthesis of α-Substituted Serines", Tetrahedron Letters, <u>36</u>, No. 23, pp. 4101-4104 (1995)</p> <p>NAGAO, et al., "Efficient Preparation of New Chiral Synthons Useful for (+)-Carbacyclin Synthesis by Utilizing Enzymatic Hydrolysis" Chemistry Letters, pp. 239-242 (1989)</p> <p>TAMAI et al., "Enzymatic Hydrolyses of the α-Symmetric Dicarboxylic Diesters Bearing a Sulfinyl Group as the Prochiral Center, Chemistry Letters, pp. 2381-2384 (1994)</p> <p>CASARRUBIO, et al., "On the Syntheses of Thiophene Analogs of Practolol and 'Reversed' Practolol", J. Heterocyclic Chem, <u>20</u>, 1557-1560 (1983)</p> <p>CHARETTE et al., "Synthesis of α, α-Disubstituted-α-Amino Acids by Double Nucleophilic Addition to Cyanohydrins", Tetrahedron Letters <u>39</u>, 5147-5150 (1998)</p>					
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				Examiner Name		
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OTHER DOCUMENTS - NON-PATENT LITERATURE DOCUMENTS						
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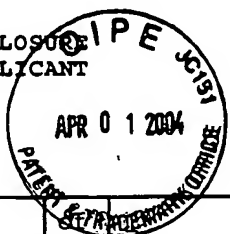
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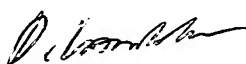


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Substitute for Form 1449A/PTO INFORMATION DISCLOSURE STATEMENT BY APPLICANT 		Application Number	10/718,858
		Filing Date	November 20, 2003
		First Named Inventor	Takahide NISHI et al.
		Group Art Unit	
		Examiner Name	
Sheet	1	Attorney Docket Number	02697CIPD/HG

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<i>PN</i>		KIUCHI et al., "Synthesis and Biological Evaluation of 2,2-Disubstituted 2-Aminoethanols: Analogues of FTY720" BIOORGANIC AND MEDICINAL CHEMISTRY LETTERS, vol. 8, 1998, pages 101-106 XP002271143	
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			3/21/05

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Substitute for Form 1449A/PTO		Application Number	10/718,058
INFORMATION DISCLOSURE STATEMENT BY APPLICANT		Filing Date	November 20, 2003
		First Named Inventor	Takahide NISHI
		Group Art Unit	1614
		Examiner Name	
Sheet	1	of	1
		Attorney Docket Number	02697CIPD/EG

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Examiner Signature	<i>D-Lanblum</i>	Date Considered	<i>3/21/05</i>
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